

# Is the Factor of 10 Still Applicable Today?

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# Disclaimer

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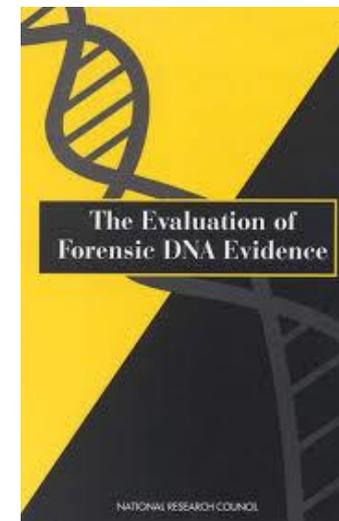
Methods

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Conclusions

# Factor of 10

NRC II, Overview, page 39:

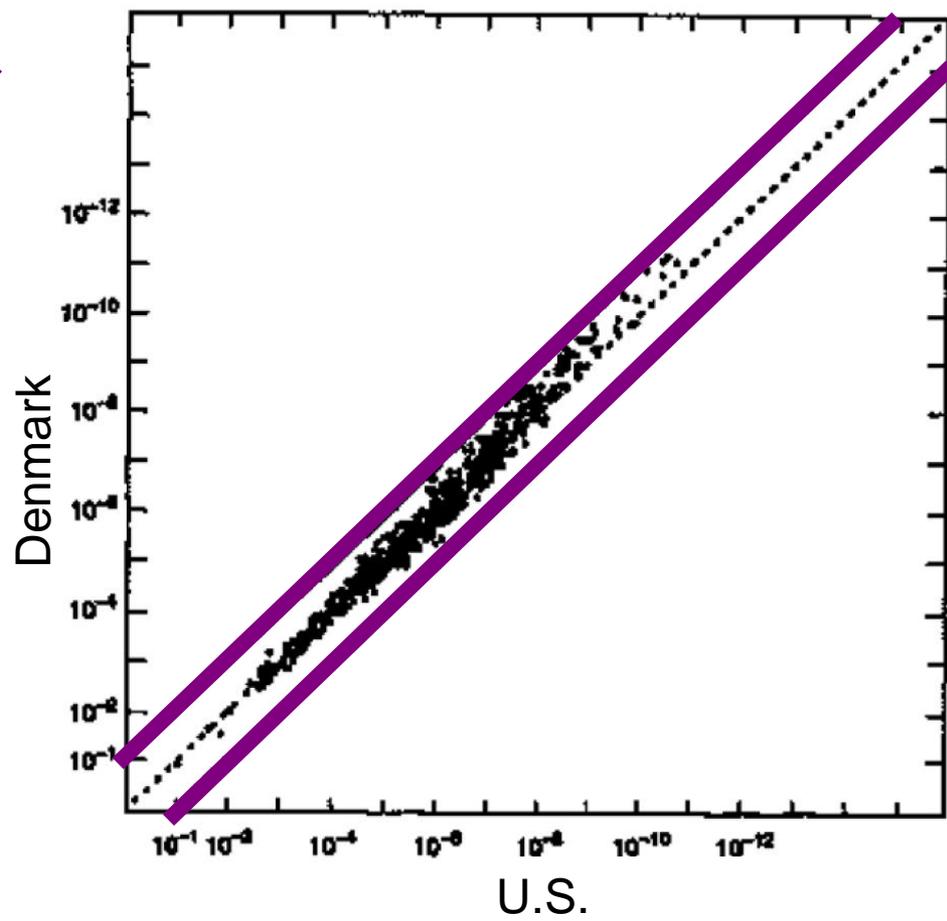
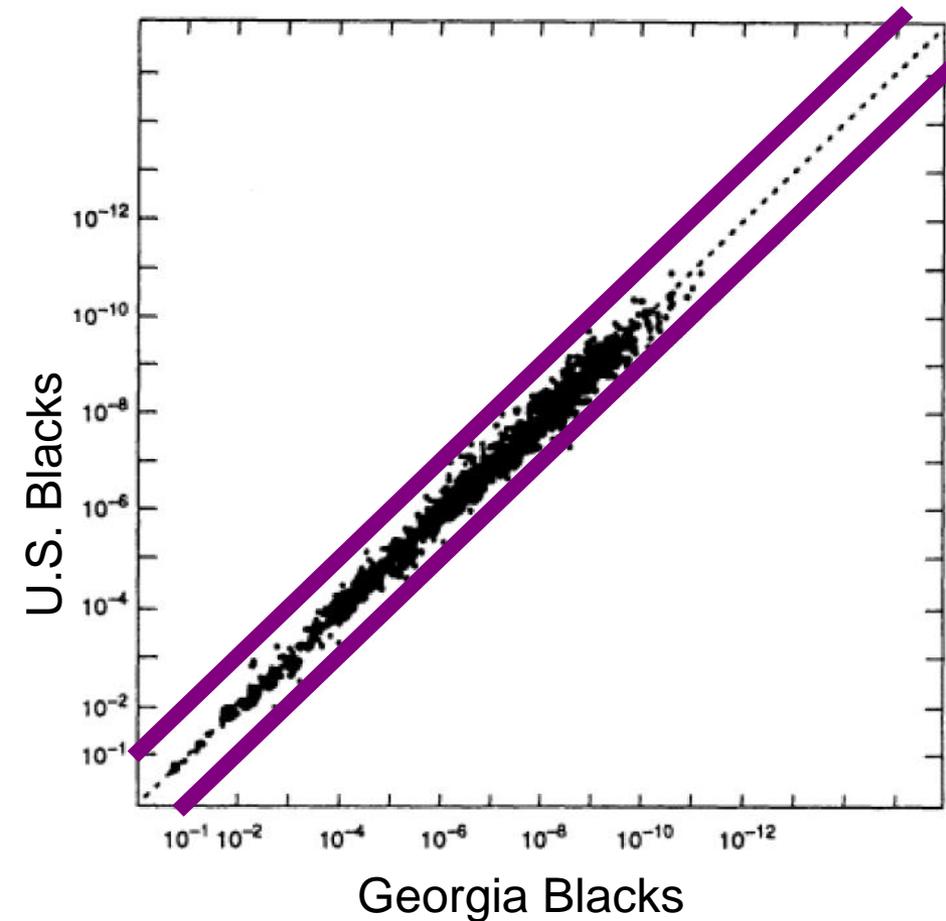
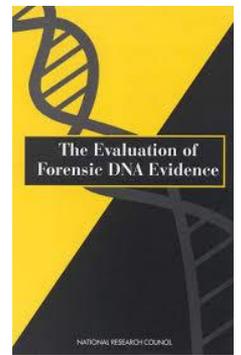


“(...) we examined empirical data from the comparison of different subpopulations and of subpopulations within the whole. The empirical studies show that the differences between the frequencies of the individual profiles estimated (...) from different adequate subpopulation databases (...) are **within a factor of about 10 of each other,**”  
*(emphasis added by me)*

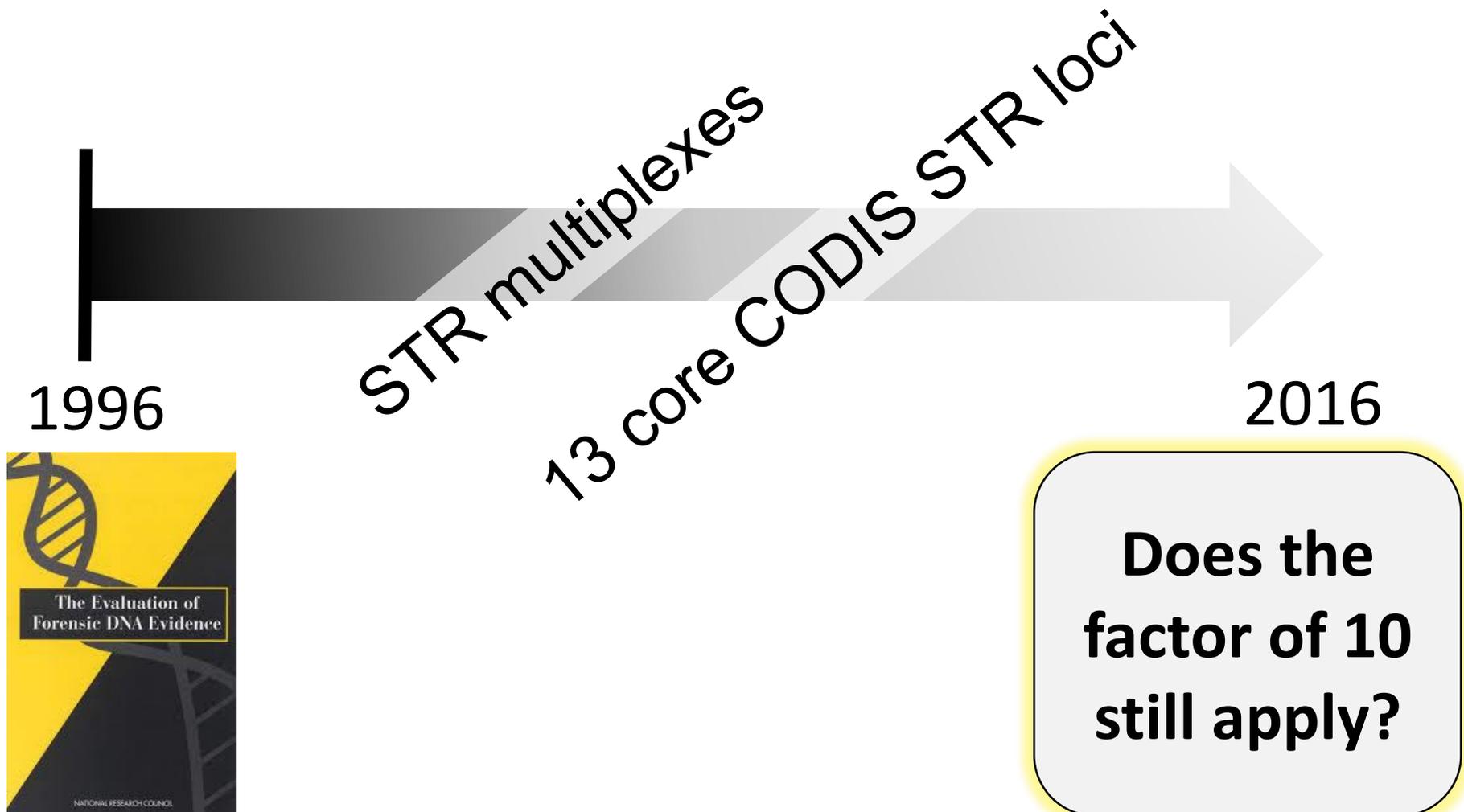
**National Research Council Committee on DNA Forensic Science.** The Evaluation of Forensic DNA Evidence. Washington, D.C.: National Academy Press; 1996.

# Factor of 10

NRC II, Statistical Issues, pages 150 and 152:



# Factor of 10



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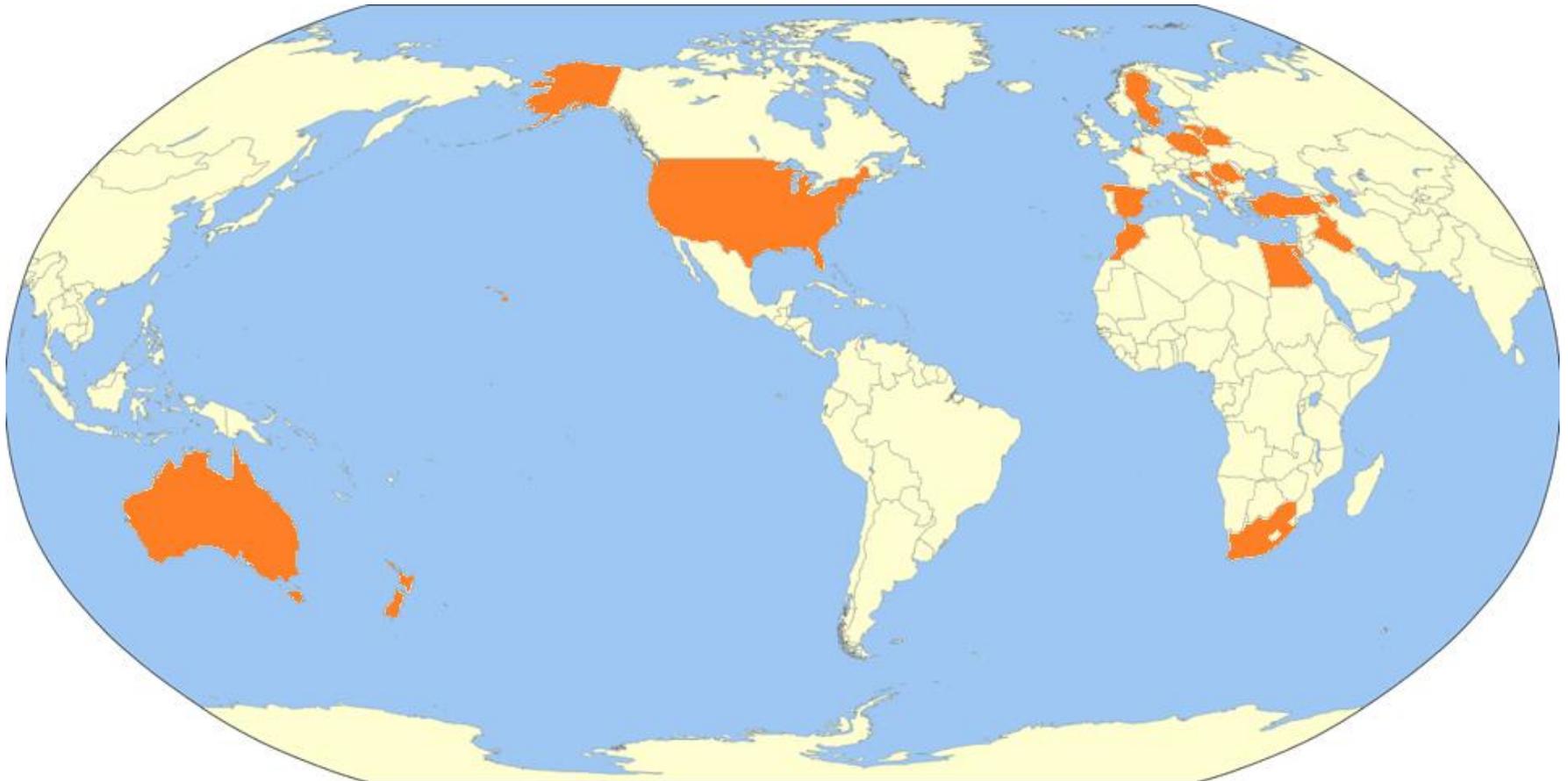
Conclusions

# Methods

AmpF $\ell$ STR $\text{\textcircled{R}}$  Identifiler $\text{\textcircled{R}}$  PCR Amplification Kit

AmpF $\ell$ STR $\text{\textcircled{R}}$  Identifiler $\text{\textcircled{R}}$  Plus PCR Amplification Kit

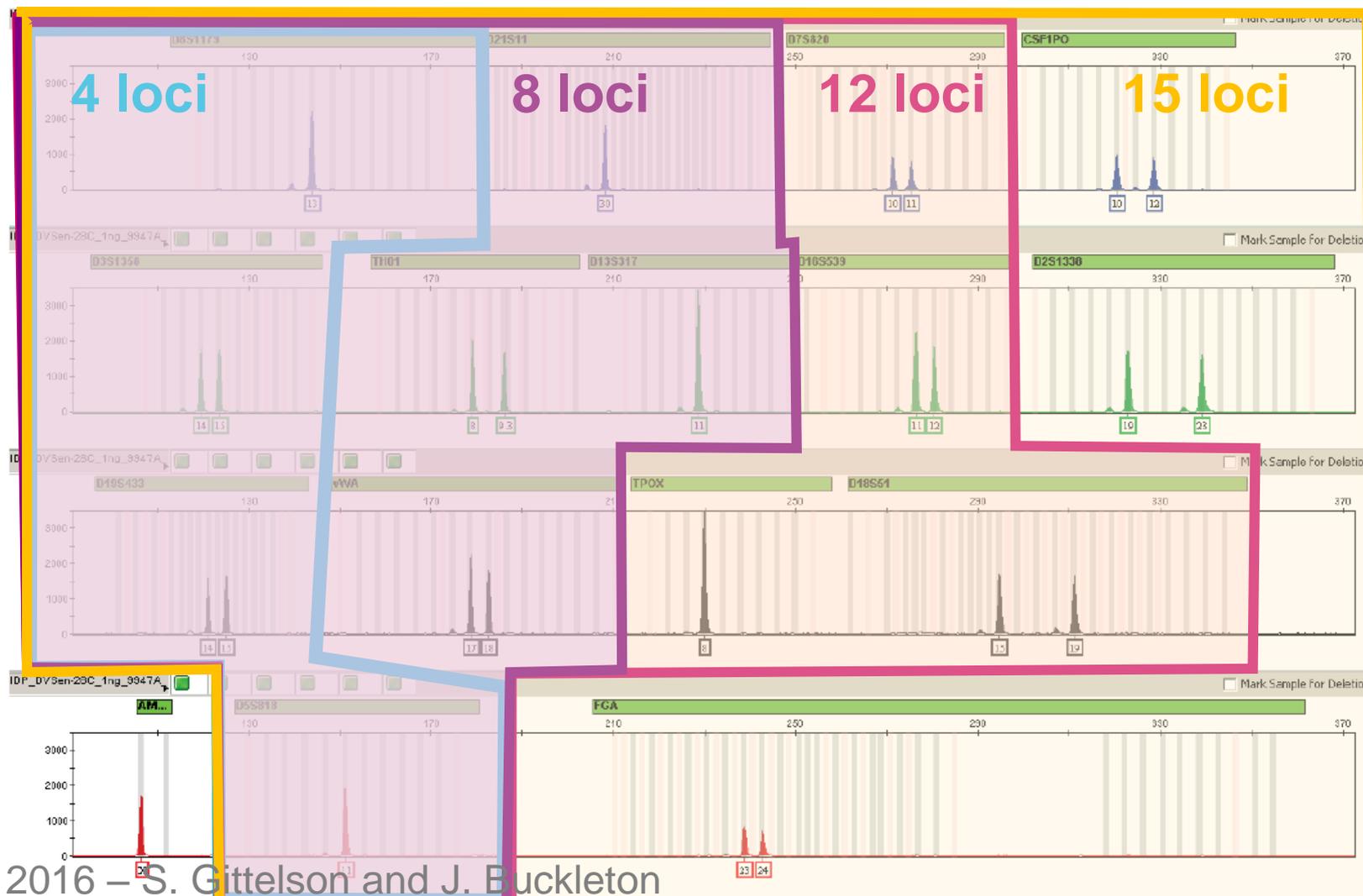
Caucasians:



# Methods

AmpF $\ell$ STR $^{\circledR}$  Identifiler $^{\circledR}$  PCR Amplification Kit

AmpF $\ell$ STR $^{\circledR}$  Identifiler $^{\circledR}$  Plus PCR Amplification Kit



# Methods

Simulations of:

4 loci



100 profiles

8 loci



100 profiles

12 loci



100 profiles

15 loci



100 profiles

= 400 profiles per subpopulation

# Profile Probability

NRC II, Recommendation 4.1.:

homozygote

heterozygote

$$Fp_i + (1 - F)p_i^2$$

$$2p_i p_j$$

→ takes into account inbreeding, but not co-ancestry

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# Results

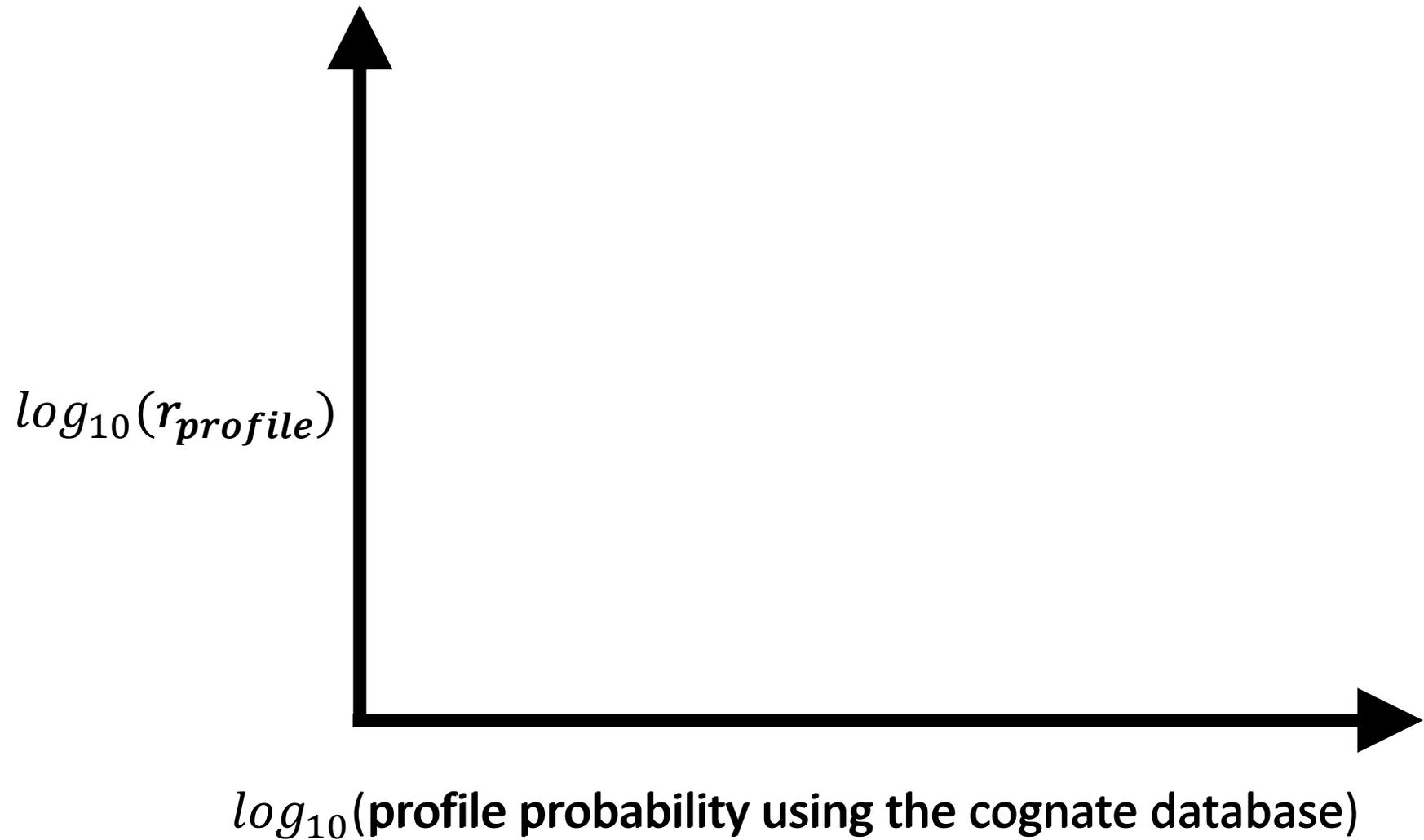
$$r_{profile} = \frac{\text{profile probability using non-cognate database}}{\text{profile probability using cognate database}}$$

if  $r_{profile} = 1$ : non-cognate profile probability is the same as the cognate profile probability

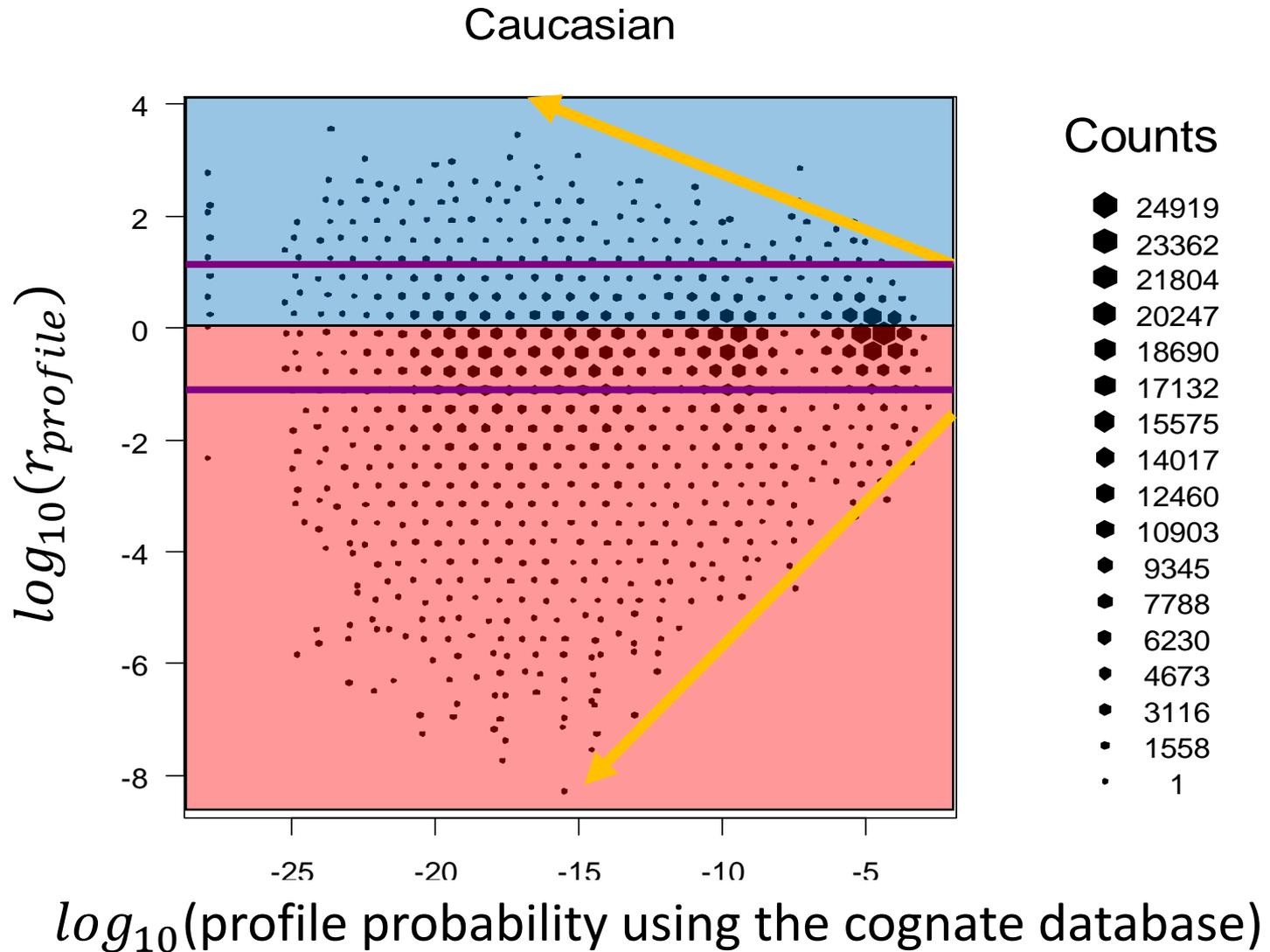
if  $r_{profile} > 1$ : the profile is more common in the non-cognate database than in the cognate database

if  $r_{profile} < 1$ : the profile is rarer in the non-cognate database than in the cognate database

# Results

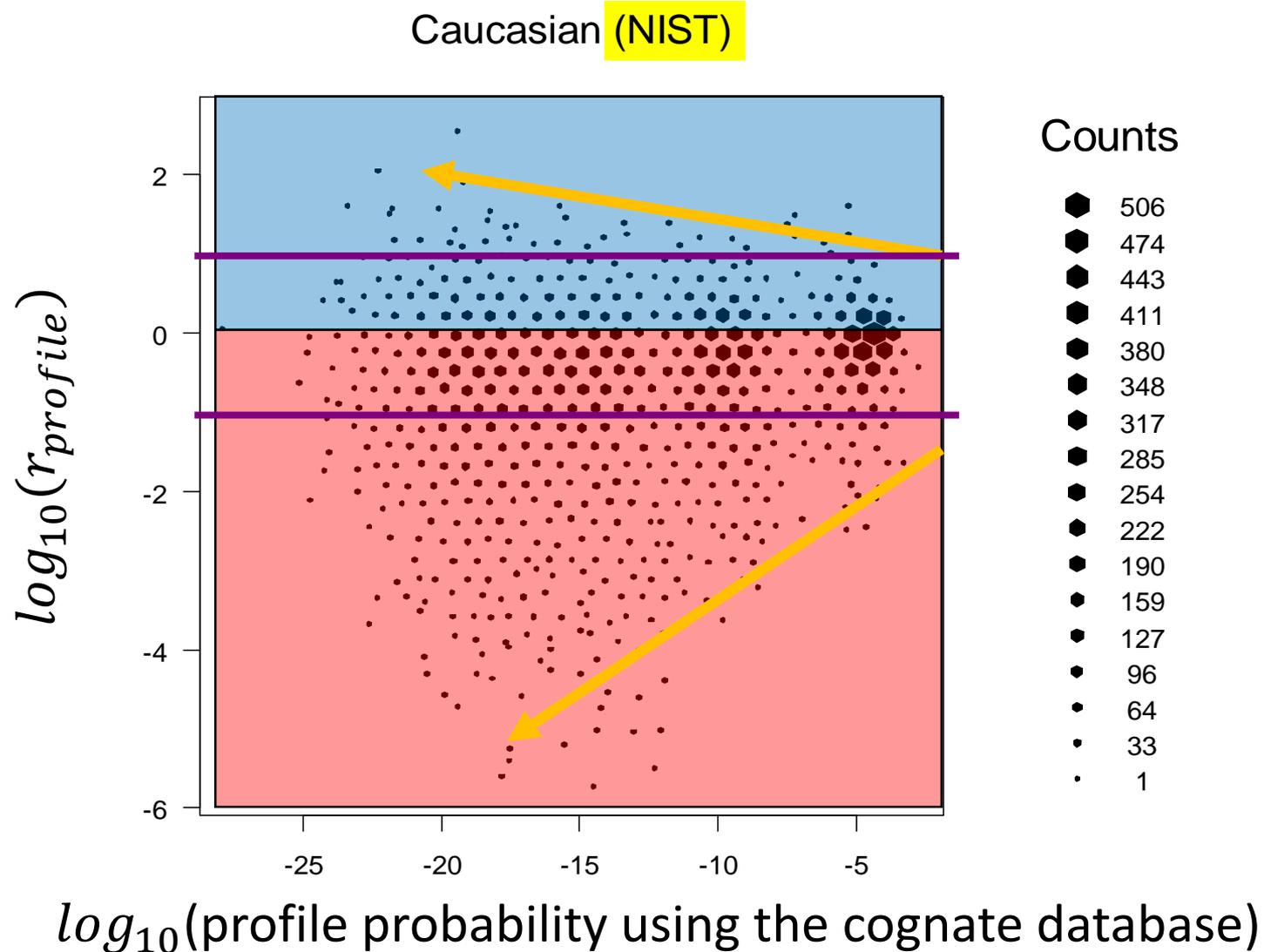


# Results



# Results

Hill CR, Duewer DL, Kline MC, Coble MD, Butler JM. US population data for 29 autosomal STR loci. *Forensic Sci. Int.: Genet.* 2014; 7: e82-e83.



# Match Probability

NRC II, Recommendation 4.2.:

homozygote

$$\frac{(2\theta + (1-\theta)p_i)(3\theta + (1-\theta)p_i)}{(1+\theta)(1+2\theta)}$$

heterozygote

$$2 \frac{(\theta + (1-\theta)p_i)(\theta + (1-\theta)p_j)}{(1+\theta)(1+2\theta)}$$

→ takes into account inbreeding and co-ancestry

**Balding DJ, Nichols RA.** DNA profile match probability calculation: how to allow for population stratification, relatedness, database selection and single bands. *Forensic Sci. Int.* 1994; 64: 125-140.

# Results

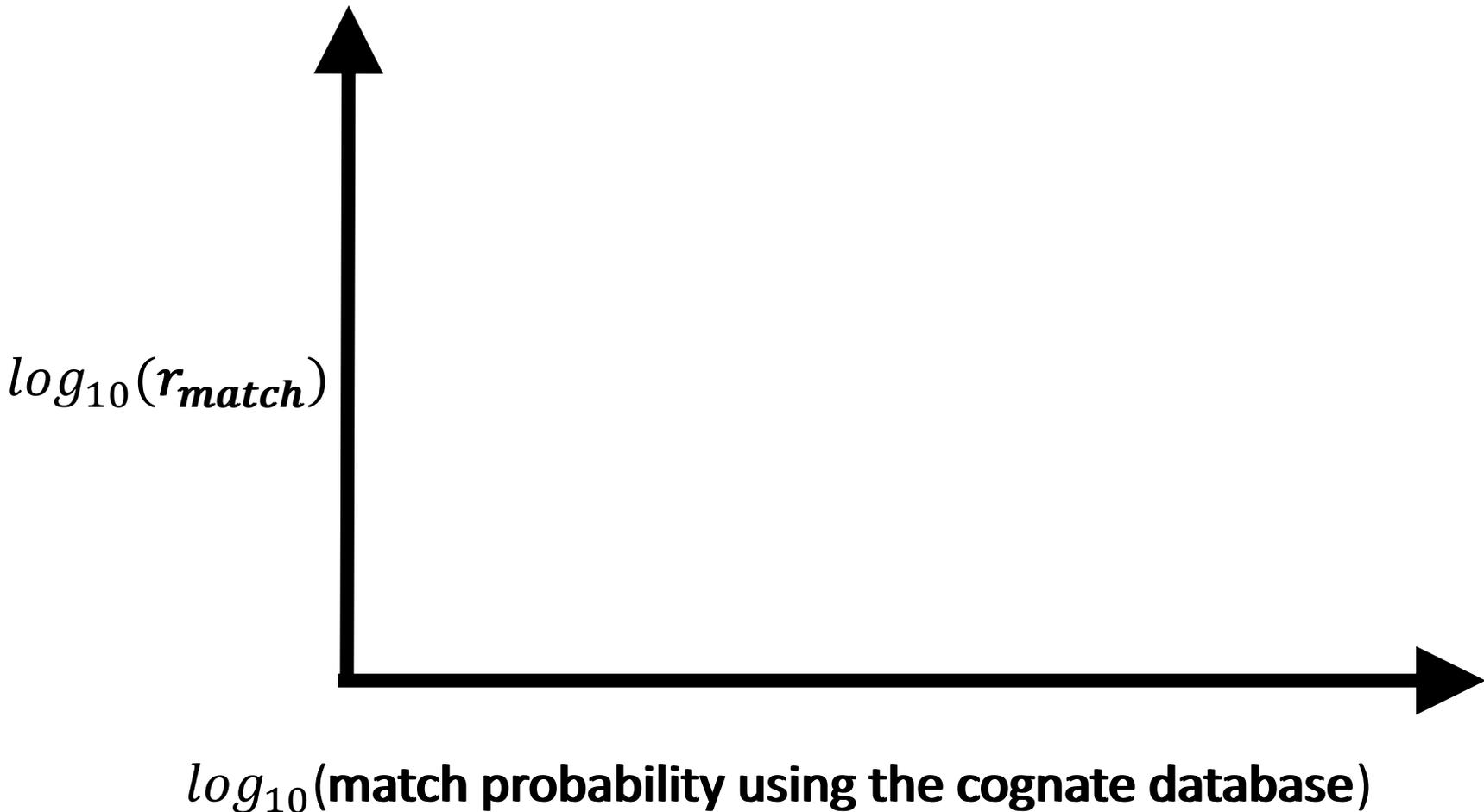
$$r_{match} = \frac{\text{match probability using non-cognate database}}{\text{match probability using cognate database}}$$

if  $r_{match} = 1$ : non-cognate match probability is the same as the cognate match probability

if  $r_{match} > 1$ : a match is more common in the non-cognate database than in the cognate database

if  $r_{match} < 1$ : a match is rarer in the non-cognate database than in the cognate database

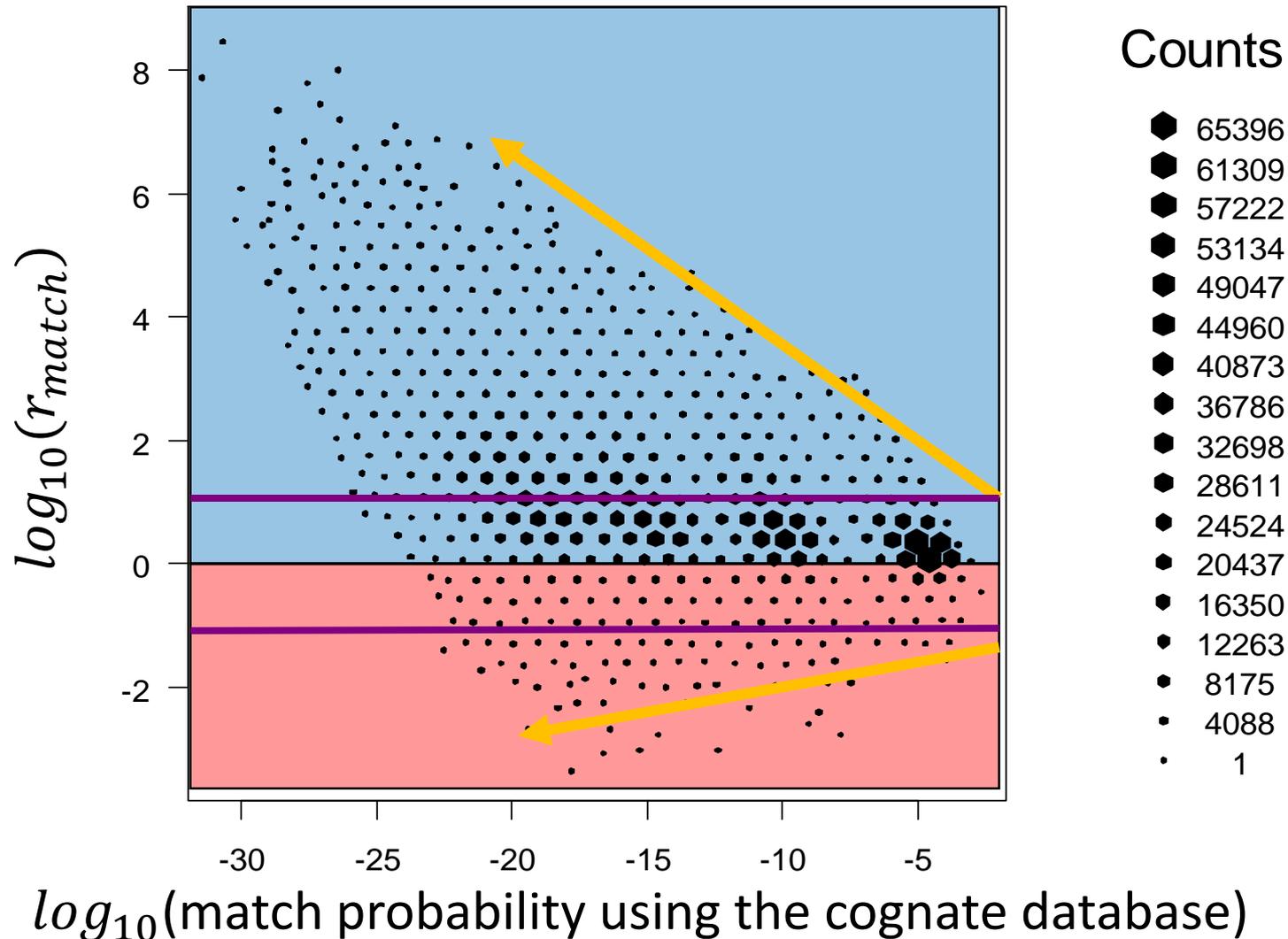
# Results



# Results

Hill CR, Duewer DL, Kline MC, Coble MD, Butler JM. US population data for 29 autosomal STR loci. *Forensic Sci. Int.: Genet.* 2014; 7: e82-e83.

Caucasian (NIST)



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# Conclusions

Variation can be greater than a factor of 10.

Non-cognate **profile probabilities** tend to be **rarer** than the cognate profile probabilities.

→ **non-conservative**

Non-cognate **match probabilities** tend to be **more common** than the cognate match probabilities.

→ **conservative**

Thank you very much for your  
attention!